A C-TERMINAL HAEM-BOUND CYTOCHROME *c*-556 FROM *AGROBACTERIUM TUMEFACIENS*, STRAIN B_2a

Amino acid composition and N-terminal sequence analysis

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1. Introduction

Comparison of amino acid sequences of soluble cytochromes *c* is a successful approach to unravel eukaryotic phylogeny [1]. For the prokaryotes, the same approach has, until now, almost exclusively been applied to the cytochromes of some phototrophic bacteria, of *Paracoccus denitrificans*, of some *Desulfovibrio* and *Pseudomonas* strains and of *Azotobacter vinelandii* [1]. Cytochromes of aerobically grown chemo-organotrophes have not been investigated [1–3]. Because of its peculiar nature, we report here on the N-terminal sequence of cytochrome *c*-556 from *Agrobacterium tumefaciens*, strain B_2a. It is a soluble protein with mol. wt ~12 000. Its purification and properties have been described [4]. In addition to our recent findings on the sequence of the haem-binding region of the protein [5], we conclude that B_2a *c*-556 is a C-terminal haem-bound cytochrome.

2. Materials and methods

The preparation of B_2a *c*-556 was electrophoretically pure and had a purity index of 1.47 [4]. Amino acid analyses were carried out with a Biocal 200 analyser after hydrolysis of the native protein with 6 N HCl at 105°C in sealed, evacuated ampoules. The N-terminal sequence of the protein was determined in a Socosi PS 110 sequencer using the 1 N quadrol programme in [6]. The thiazolinones were converted with 0.2 ml 1 N HCl at 80°C. After 3 extractions with 0.7 ml ethylacetate, the PTH-amino acids in the organic phase were identified on Kieselgel thin-layer plates (Merch 5715, 20 × 20 cm). One-half of the aqueous phase was assayed for PTH-arginine [7], the other half for PTH-histidine [8], both on paper. Three tryptic peptides which cover the N-terminal region of the cytochrome were obtained by digestion of 2 μmol dehaemed protein. Their sequence was determined by manual dansyl-Edman degradation.

Abbreviations: PTH, phenyl thiohydantoin amino acids; B_2a *c*-556, cytochrome *c*-556 from *A. tumefaciens*, strain B_2a

3. Results and discussion

The amino acid composition of B_2a *c*-556 is in table 1. Since –Cys–X–Y–Cys–His– is the haem-binding sequence in most cytochromes *c* (except for –Ala–X–Y–Cys–His– in two eukaryotic cytochromes [1]), the occurrence of 2 cysteines and 1 histidine indicates that B_2a *c*-556 is a monohaem cytochrome. The mol. wt 12 540 calculated from the presumed composition is in good agreement with the value obtained by SDS–gel electrophoresis [4].

Figure 1 shows the N-terminal sequence of B_2a *c*-556. The first 16 residues were identified with the native protein in the sequenator. No further residues could be identified, probably because of a nearly complete cyclisation of Gln-13 to pyrrolidone–car-

Table 1
Amino acid composition of *A. tumefaciens* B_2a *c*-556

Amino acid	Residues/mol protein after hydrolysis at		Nearest integer
	24 h	96 h	
Asp	9.8	8.6	10
Thr[a]	9.8	9.2	10
Ser[a]	2.9	2.9	3
Glu	12.0	12.4	12
Pro	4.5	4.5	5
Gly	13.3	12.9	13
Ala	20.8	20.9	21
Val	4.4	4.5	4
Met	1.8	3.2	3
Ile	4.1	4.1	4
Leu	6.0	5.8	6
Phe	5.0	4.9	5
Lys	10.8	12.0	12
His	0.7	1.1	1
Arg	2.0	2.0	2
Tyr	0.7	1.0	1
Cys[b]	2	2	2
Trp			1[c]

[a] Corrected values for decomposition during acid hydrolysis
[b] Determined as cysteic acid
[c] Estimated from the purity of the native cytochrome

boxylic acid; the repetitive yield for the PTH of Glu-8 and Gly-9 was found to be as high as 95%. Sequence information beyond residue 16 could nevertheless be obtained by analysing those peptides of a tryptic protein digest which fit the N-terminal sequence found with the sequenator. The peptides T3e and T3d3 (fig.1), constitute the region 1 → 12 of the protein. T2b3 is the only other tryptic peptide with N-terminal Glu or Gln (determined as dansyl-Glu). Its subsequent sequence –Ile–Glu–Gly– is identical with the last 3 amino acids determined with the sequenator. T2b3 is thus the neighbouring peptide of T3d3, covering the region 13 → 26.

It is clear from our data that no cysteines or histidine occur within the 26 N-terminal positions of B_2a *c*-556. In most N-terminal haem-bound cytochromes (they are grouped under class I according to the classification in [2]) the first cysteine of the invariant haem-binding sequence –Cys–X–Y–Cys–His– is at or before position 14 [1]. The most remote position in eukaryotic cytochromes of class I is at position 25 (*Tetrahymena pyriformis* [9]), in prokaryotic cytochromes of the same class it is at position 23 (*Prosthecochloris aestuarii* [10,11]). We therefore conclude that B_2a *c*-556 is not an N-terminal haem-bound cytochrome.

Within the range of the presently studied structures, the only other alternative is that B_2a *c*-556 is a member of class II (A classification [2]). This class contains high- (cytochromes *c'*) and low-spin cytochromes with the haem group bound at the C-terminal part of the polypeptide chain. Our conclusion is strengthened by the sequence homology of the haem region of B_2a *c*-556 with the C-terminal region of the following class II cytochromes examined: cytochrome *c*-556 (low-spin) from *Rhodopseudomonas palustris*; the cytochromes *c'* from *Rps. capsulata, Rps. palustris, Chromatium vinosum, Rhodospirillum rubrum* from one *Alcaligenes* strain, NCIB 11015 (fig.2b). The primary structure of the latter 2 cytochromes *c'* are the only ones completely elucidated so far. The similarity between the haem regions from the latter 2 proteins and B_2a *c*-556, reported [5], was 35%. As a result of the present work, we also aligned the N-terminal region of the *Agrobacterium* cytochrome with the corresponding region of both cytochromes *c'* (fig.2a). The similarity is clearly lower than at the C-terminus: it is 27% with *Rhs. rubrum* cytochrome *c'* and 15% with the *Alcaligenes* protein; it is also 27% between the 2 cytochromes *c'*. Three residues seem to be evolutionary invariant: Val-4, Arg-7 and Lys-26. All these data support the conclusion that

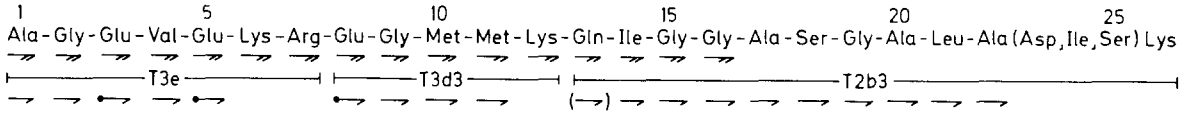

Fig.1. N-terminal amino acid sequence of *A. tumefaciens* B_2a *c*-556. T3e, T3d3 and T2b3: tryptic peptides. (—//) Residues identified with the sequenator; (—$_7$) residues identified as the dansyl amino acid; (•—$_7$) residues identified as the dansyl derivate of the acid but final acid assignment deduced from the mobility of the peptide; ((—$_7$)) acid or amide which cannot be deduced from the mobility alone.

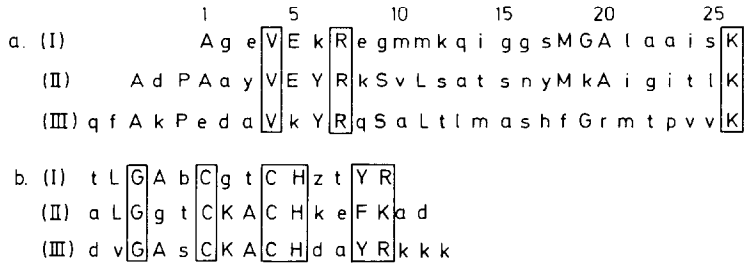

Fig.2. Comparison of the N-terminus (a) and the C-terminal haem region (b) of *A. tumefaciens*, cytochrome *c*-556 (I) and the cytochromes *c'* from *Rhs. rubrum* (II) and *Alcaligenes* NCIB 11015 (III). The one-letter code follows the recommendations in [14]. Capital letters are used when a residue in any one sequence is identical to a residue in another sequence. Boxed residues, including the substitution Tyr ↔ Phe and Arg ↔ Lys at the C-terminus, are common to the 3 cytochromes. The boxed residues in (b) are also common to the 4 other cytochromes of class II mentioned in the text.

A. tumefaciens B_2a *c*-556 is a class II cytochrome *c*.

As a point of structural and phylogenetic interest, it is remarkable that the first cytochrome *c* studied from an aerobically grown chemo-organotrophe belongs to a cytochrome class of which the members, with exception of the cytochromes *c'* from *Azotobacter vinelandii* [12], a halotolerant 'micrococcus' [3,13] and one strain of *Alcaligenes*, are found exclusively in photosynthetic bacteria. The complete sequence determination of B_2a *c*-556 is in progress.

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